Study Director General Toxicology

Job Offer

AnaPath Research

AnaPath Research is a CRO located in Barcelona, with extensive experience in carrying out preclinical trials for pharmaceutical laboratories, the chemical industry and other research organizations. In our more than 30 years of activity we have worked with the main pharmaceutical industries being part of different multinationals (RCC, Harlan and Envigo). In November 2019, AnaPath Services acquired the company and re-founded it as AnaPath Research, thus undertaking together a new project of scientific quality and close contact with new and old sponsors.

With a multidisciplinary team of scientific experts, AnaPath Research covers most fields of preclinical pharmaceutical development and chemical and food safety.

Department

The General Toxicology Department performs a full range of studies to support the pharmaceutical, chemical, agrochemical and veterinary industries.

Our team consists of senior scientists with an average of 20 years of experience in toxicology, pharmacokinetics and safety pharmacology, and wide knowledge of the current regulatory guidelines.

We offer scientific advice on study types, study design and choice of dose levels before starting with first-in-human trials.

As a GLP-certified and AAALAC-accredited center, we fully embrace the 3Rs of animal welfare. We are strongly convinced that our animal welfare policies have a positive impact on our studies

Position

We are looking for a Study Director for general toxicology, safety assessment, reprotoxicology, pharmacokinetics, and veterinary regulatory studies.

AnaPath is accredited to work with pharmacological, chemical, food additives and phytosanitary products.

The team of the Toxicology Department, like the rest of the company, is committed to animal welfare and possess an extensive knowledge in the preclinical development of drugs.

Responsibilities

Act as Study Director of the General Toxicology Department.

Functions will include study plan (and amendment) design and approval in accordance with internal procedures and, where appropriate, in compliance with international regulations; study coordination; design, development and approval of the study work schedule; interpretation of the results; writing of the discussion and conclusion for the final report. Tasks will be carried out in accordance with the instructions received from the manager and the sponsor (evaluating the procedures proposed by the sponsor in addition to providing relevant solutions when necessary) and in agreement with the rest of the participating staff.

As Study Director he/she will be the person responsible for the studies assigned to him/her by the managers, i.e., before the start of the studies, he/she should check and confirm that the staff and resources required for the proper management and conduct of the studies are available.

Requirements

Life sciences degree, Scientific PhD or MSc.

Expertise: minimum of 5 years in general toxicology/safety pharmacology studies and/or veterinarian TAS studies

Previous experience in a research laboratory is an asset

Qualification to act as investigator (d) (according to EU Directive 63/2010, RD 53/2013 (Royal Decree) and ECC/566/2015 (Official State Gazette)).

Good knowledge of good laboratory practice (GLP) standards.

Good knowledge of Animal Welfare regulations.

Knowledge of regulatory toxicology is an asset

Good communication skills

Advanced English level

Work permit for Spain

Terms of Employment

Indefinite contract

Full-time

Morning work shift

Contact

rrhh@anapathresearch.com